Listing of Claims:

1 (currently amended): A non-reactive pressure sensitive adhesive composition comprising an acrylic polymer and a therapeutic agent, wherein the acrylic polymer

(i) is prepared from monomers selected from the group consisting of alkyl acrylate monomers, alkyl methacrylate monomers, polymerizable non-cyclic nitrogen-containing monomers and mixtures thereof.

wherein said alkyl acrylate monomers and alkyl methacrylate monomers have up to about 18 carbon atoms in the alkyl group, and

wherein said polymerizable non-cyclic nitrogen-containing monomers are selected from the group consisting of t-octyl acrylamide, dimethyl acrylamide, diacetone acrylamide, t-butyl acrylamide, i-propyl acrylamide, N-phenyl acrylamide, vinylacetamides, nitriles, and mixtures thereof, and

wherein said alkyl acrylate monomers and/or alkyl methacrylate monomers are present in the acrylic polymer in amounts of from about 50 to about 98%, based on a dry weight basis of the total monomer weight of the acrylic polymer, and said polymerizable noncyclic nitrogen-containing monomers are present in the acrylic polymer in amounts of from about 2 to about 50%, based on a dry weight basis of the total monomer weight of the acrylic polymer,

- (ii) lacks functional groups containing reactive hydrogen mojeties and
- (iii) contains no post-polymerization chemical crosslinking[,] and wherein the therapeutic agent is a non-salt agent.

Atty. Docket No. N-1893.TDM-US Application No.: 09/955,644 Office Action Dated: October 21, 2010

2 canceled

3 (previously presented): The adhesive of claim 1 wherein the acrylic polymer is prepared using a

nitrile, which nitrile is methacrylonitrile or 2-cyanoethylacrylate.

4 (original): The adhesive of claim 1 which has a Tg of less than about 10°C.

5 (previously presented): The adhesive of claim 4 wherein the acrylic polymer is prepared using an

alkyl acrylate monomer, which alkyl acrylate monomer is 2-ethylhexyl acrylate and/or n-butyl

acrylate.

6 (previously presented): The adhesive of claim 5 wherein the acrylic polymer is prepared using an

N-substituted acrylamide monomer and/or an N-substituted methacrylamide monomer.

7 (original): The adhesive of claim 6 wherein N-substituted acrylamide is t-octyl acrylamide.

8 canceled

9 (previously presented): The adhesive of claim 1 wherein the therapeutic agent is a

pharmacologically active agent.

10 (previously presented): A transfermal drug delivery system comprising the adhesive of claim 1.

11 (previously presented): The transdermal drug delivery system of claim 22 wherein the adhesive

serves as a carrier for the therapeutic agent.

12 (currently amended): A transdermal drug delivery system comprising a non-reactive pressure

sensitive adhesive layer and a backing layer, wherein said adhesive layer comprises

(a) an acrylic polymer, wherein said acrylic polymer

(i) is prepared from monomers selected from the group consisting of alkyl acrylate

monomers, alkyl methacrylate monomers, polymerizable non-cyclic nitrogen-containing

monomers and mixtures thereof,

wherein said alkyl acrylate monomers and alkyl methacrylate monomers have up to

about 18 carbon atoms in the alkyl group, and

wherein said polymerizable non-cyclic nitrogen-containing monomers are selected

from the group consisting of t-octyl acrylamide, dimethyl acrylamide, diacetone acrylamide,

t-butyl acrylamide, i-propyl acrylamide, N-phenyl acrylamide, vinylacetamides, nitriles, and

mixtures thereof, and

wherein said alkyl acrylate monomers and/or alkyl methacrylate monomers are

present in the acrylic polymer in amounts of from about 50 to about 98%, based on a dry

weight basis of the total monomer weight of the acrylic polymer, and said polymerizable non-

cyclic nitrogen-containing monomers are present in the acrylic polymer in amounts of from

about 2 to about 50%, based on a dry weight basis of the total monomer weight of the acrylic

Atty. Docket No. N-1893.TDM-US Application No.: 09/955,644 Office Action Dated: October 21, 2010

polymer,

(ii) lacks functional groups containing reactive hydrogen moieties and

(iii) contains no post-polymerization chemical crosslinking

and

(b) a therapeutic agent, which is fentanyl.

13 (original): The transdermal drug delivery system of claim 12 further comprising a release layer.

14 (previously presented): A method of administering a therapeutic agent to a patient comprising applying to a body surface of a patient the transfermal drug delivery system of claim 12.

15 (previously presented): The adhesive of claim 9 wherein the pharmacologically active agent is fentanyl.

16 (previously presented): A transdermal drug delivery system comprising the adhesive of claim 15.

17 (previously presented): The method of claim 14 wherein the therapeutic agent is fentanyl.

18 (previously presented): The adhesive of claim 1 comprising an acrylic polymer prepared from 2ethylhexyl acrylate, methyl acrylate and an N-substituted acrylamide monomer.

19 (previously presented): The adhesive of claim 18 wherein the nitrogen-containing acrylamide

monomer is t-octyl acrylamide.

20 canceled

21 (previously presented): The transdermal drug delivery system of claim 12 comprising an acrylic

polymer prepared from 2-ethylhexyl acrylate, methyl acrylate and t-octyl acrylamide.

22 (currently amended): A transdermal drug delivery system comprising a non-reactive pressure

sensitive adhesive and a therapeutic agent, wherein

said adhesive comprises an acrylic polymer, which acrylic polymer

(i) is prepared from monomers selected from the group consisting of alkyl acrylate

monomers, alkyl methacrylate monomers, polymerizable non-cyclic nitrogen-containing

monomers and mixtures thereof,

wherein said alkyl acrylate monomers and alkyl methacrylate monomers have up to

about 18 carbon atoms in the alkyl group, and

wherein said polymerizable non-cyclic nitrogen-containing monomers are selected

from the group consisting of t-octyl acrylamide, dimethyl acrylamide, diacetone acrylamide,

t-butyl acrylamide, i-propyl acrylamide, N-phenyl acrylamide, vinylacetamides, nitriles, and

mixtures thereof, and

wherein said alkyl acrylate monomers and/or alkyl methacrylate monomers are

present in the acrylic polymer in amounts of from about 50 to about 98%, based on a dry

weight basis of the total monomer weight of the acrylic polymer, and said polymerizable non-

Atty. Docket No. N-1893.TDM-US Application No.: 09/955,644

Office Action Dated: October 21, 2010

cyclic nitrogen-containing monomers are present in the acrylic polymer in amounts of from

about 2 to about 50%, based on a dry weight basis of the total monomer weight of the acrylic

polymer,

(ii) lacks functional groups containing reactive hydrogen moieties and

(iii) contains no post-polymerization chemical crosslinking[,]

and wherein the therapeutic agent is a non-salt agent.

23 (previously presented): The adhesive of claim 18 wherein the monomer composition of the acrylic

polymer is 45 % by weight 2-ethylhexyl acrylate, 35 % by weight methyl acrylate and 20 % by

weight of an N-substituted acrylamide monomer.